PA31A-0815 Producing Science to Inform Policy on Hemispheric Transport of Air Pollution

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The Task Force on Hemispheric Transport of Air Pollution



Task Force on Hemispheric Transport of Air Pollution

Chairs: Terry Keating (United States) and André Zuber (European Community)

Mission:

To develop a fuller understanding of the hemispheric transport of air pollution to inform future negotiations under the Convention on Long Range Transboundary Air Pollution (CLRTAP).

Background: Evidence from both observational and modeling studies indicates that pollutant emissions from one continent (source) influence air quality over other continents (receptors). A lack of community-wide consistency in defining source-receptor regions and in reporting air quality metrics, however, hinders assessment of the policy implications of intercontinental transport.

Products:

2007 Interim Report to inform the review of the CLRTAP Gothenburg Protocol to abate acidification, eutrophication, and tropospheric ozone.

2009 Assessment Report to inform the CLRTAP about hemispheric air pollution and source-receptor relationships for ozone, aerosols, mercury, and POPs.

Challenge:

Balancing the limited time and resources of individual research groups available for a community-wide effort with the needs of the policy community.

Opportunity: Framework to move information from the science to the policy community fosters a rapid exchange of ideas, helping to identify research priorities and gain insights through collaboration.

Approach:

- 1. Define a set of focused, policy-relevant questions.
- 2. Engage the scientific community in designing a set of experiments to address these questions.
- 3. Translate science results into metrics that are directly relevant to air quality policy goals.
- 4. Maintain an open dialogue among participants via several communication avenues.

A Multi-model Assessment

Objectives:

Quantify source-receptor relationships for HTAP regions (see top right) and assess uncertainties in these estimates.

Methods:

- 1. Encourage participation of groups with global 3D tropospheric chemistry models (require horizontal resolution of 4°x5° or finer).
- 2. Specify base case simulation: 2001 meteorology; methane set to a uniform value of 1760 ppb; each group uses its best estimate for NO., NMVOC, CO emissions.
- 3. Conduct sensitivity simulations: 20% reductions of anthropogenic NO., NMVOC, and CO individually from each HTAP region, and a 20% reduction of CH₄ abundances to 1408 ppb (13 total simulations).
- 4. Require model output to comply with CF conventions (a version of the Climate Model Output Rewriter is available for this task).

Encouraging Dialogue: TF HTAP Communication Avenues



To join the TF HTAP listsery, click or send a blank email to subscribe-tfhtap@lists.epa.gov

Meetings and Workshops

	l' di po	Purpose		
JUN 1-3, 2005 First TF Mee	ting Devel	Develop a work plan		
JAN 30-31, Modeling W		Organize multi-model assessment of HTAP and draft outline of 2007 interim report		
JUN 6-8, 2006 Second TF I	metha	Review science of mercury, POPs, and methane as an ozone precursor; agree on 2007 work plan		
OCT 18-20, Emissions I 2006 and Future I Workshop		ate current emission inventories and projections	Beijing, China	
JAN 24-26, 2007 Integrated Observations Workshop		Assess state of current observational networks; discuss how to fill gaps and produce common data bases		
JAN 27, 2007 Model Inter- comparison		ss initial analyses and priorities for interim report	Geneva, Switzerland	
MAY 30 – JUN 1, 2007 Third TF Me		ot the 2007 interim report; agree on workplan	London, U.K.	

Assessment of Hemispheric Ozone Transport: Example Analysis of Results From One Model

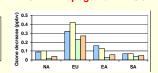
Response of seasonal mean surface O₂ over the HTAP receptor regions to 20% reductions in:

Anthropogenic NO_x

			Source	region		
듣		NA	EU	EA	SA	■ DJF
egion	NA	-1.41	-0.06	-0.09	-0.03	□ MAM
- 10	EU	-0.02	-1.31	-0.05	-0.02	□JJA
eptor	EA	-0.14	-0.15	-1.05	-0.09	SON
ĕ	SΔ	-0.13	-0.16	-0.00	-1 22	

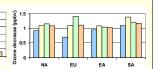
The table shows the change in seasonal mean O₃ (ppbv) resulting from a 20% reduction of anthropogenic NO. emissions in the source regions for the season of maximum impact. Table cells with equivalent impacts in multiple seasons are colored white.

EU Anthropogenic NMVOC



0.25 0.2 0.15 0.05

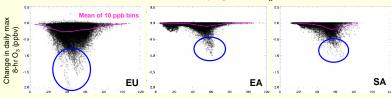
EU Anthropogenic CO



Methane

- Intercontinental influence varies by season and region
- Maximum intercontinental influence does not always occur in season with highest base case seasonal mean O₃ in the receptor region (highest base case O₃ occurs in summer for EU; spring for EA and NA; winter for SA)
- In spring, European NO₂, CO, and NMVOC contribute similarly to O₃ over NA and EA; NO₂ reductions are more effective over SA
- A 20% decrease in CH₄ lowers surface O₃ by ~1 ppbv in all HTAP regions, with EU subject to the largest seasonal variation

Intercontinental impacts of a 20% decrease in North American anthropogenic NO., in summer



Each point represents one daily maximum 8-hour average O₃ value in one model grid cell from June 1 to August 31

Base case daily max 8-hr average O₃ (ppbv)

Intercontinental influence is typically largest near the middle of the overall O₃ distribution and smaller under the cleanest and most polluted conditions

Next steps

- Distribute a visualization tool for modeling participants to examine and error-check results
- ·Analyze results from all participating models, plus new simulations (to be submitted by January 15, 2007) with NMVOC, CO, and all O, precursors simultaneously reduced from all HTAP regions
- · Use cross-model range as an estimate of uncertainty
- · Increase focus on metrics directly relevant to policy (e.g. 8-hr daily maximum O₃)
- · Incorporate results into 2007 Interim Report

All results shown here are from the GFDL MOZART-2 model

Organizations Contributing to The HTAP Modeling Effort

AMDAL/CRESS York University ICG-II, Forschungszentrum Jülich Laboratoire de Météorologie Dynamique Service d'Aéronomie, CNRS University of Maryland Baltimore County

CIEMAT CNR-IIA **ECPL-UOC** EMEP/MSC-E EMEP/MSC-W

Environment Canada NOAA/GEDI **Harvard University** IIA-CNR Italian Research Council

JRC-IES Lamar University LLNL LSCE-CEA NASA GISS

NCAR NERI UK Met Office University of Cambridge University of Edinburgh University of Oslo

NASA GSFC

We are especially grateful to Kees Cuvelier (JRC), Christiane Textor (CNRS), Philippe Thunis (JRC), Michael Schulz (LSCE) for maintaining the model intercomparison web resources, developing and documenting the HTAP-CMOR package, and processing the submitted model results.